In the Claims:

1. (currently amended) Process A process for the production of producing a compound or reaction product of the following formula (I)

[[by]] comprising the steps of converting a compound of formula (II) with 2-pyrrolidone, characterized in that an excess of wherein 2-pyrrolidone is used[[,]] in excess relative to said compound (II)[[,]] to form a reaction mixture.

- 2. (currently amended) Process The process according to claim 1, characterized in that in a subsequent further comprising the step of crystallizing said [[the]] reaction product (I) to isolate said reaction product (I) is isolated directly from [[the]] said reaction mixture by crystallisation.
- 3. (currently amended) Process The process according to claim 1, wherein said process comprises using 2-pyrrolidone in an amount of or 2, characterized in that 1.5 to 5 mol, preferably 2 to 4 mol, especially preferably 2.5 to 3.5 mol, of 2 pyrrolidone are used, each value relative to the amount of compound (II).
- 4. (currently amended) <u>Process The process</u> according to <u>any one of the preceding</u> <u>claims, characterized in that the claim 1, wherein said process comprises performing said conversion is performed at a temperature of 50 to 200 °C.</u>
- 5. (currently amended) Process The process according to claim 4, characterized in that wherein said process comprises the step temperature of the reaction mixture is initially heated heating said reaction mixture to an initial temperature of 70 to 130 °C, preferably to 80 to 120 °C, and subsequently heating said reaction mixture to a subsequent temperature of 140 to 200 °C, preferably 150 to 190 °C.
- 6. (currently amended) Process The process according to claim 5, characterized in that the wherein said process comprises the step of maintaining said initial temperature is maintained for a period of 0.5 to 2 hours h, preferably 1 to 1.5 h, and maintaining [[the]] said subsequent temperature is maintained for a period of 1 to 8 hours h, preferably from 2 to 5 h.
- 7. (currently amended) Process The process according to claim 6, characterized in that the further comprising the steps of:

cooling said reaction mixture;

seeding said reaction mixture, [[is]] after cooling, seeded with seed crystals of compound (I)[[,]]; and is maintained

maintaining said reaction mixture seeded with said seed crystals at room temperature, preferably at least 25 °C, for a period of 24 [[h]] hours to 7 days, preferably 50 to 100 h, to enable crystallisation.

- 8. (currently amended) Process The process according to claim 7, characterized in that the wherein said crystallisation is carried through at a temperature of 30 to 70 °C, preferably at 40 to 60 °C.
- 9. (currently amended) Process A process for the production of a compound of the following formula (III), comprising the following steps:

- (A) Preparing <u>said</u> compound (I) according to a process according to any one of claims claim 1 [[to 8]];
- (B) <u>performing a reduction reaction, giving to provide said compound (III) in salt form; and</u>
- (C) liberating compound (III) from the salt.
- 10. (currently amended) Process The process according to claim 9, characterized in that wherein said process comprises using a reaction product isolated by crystallisation according to any one of claims claim 1 to 8 is used as compound (I).
- 11. (currently amended) Process The process according to claim 9, comprising the step of performing said or 10, characterized in that the reduction reaction (step B) is performed in the presence of zinc and acid.
- 12. (currently amended) Process The process according to claim 11, eharacterized in that comprising the step of initially dissolving said compound (I), preferably in erystallised form, is initially dissolved in glacial acetic acid and [[that]] subsequently adding zinc and hydrochloric acid are added to said compound (I) dissolved in glacial acetic acid.
- 13. (currently amended) <u>Process The process</u> according to claim 12, characterized in that the comprising the step of performing said reduction reaction is performed in the

presence of aqueous sulfuric acid and zinc dust.

- 14. (currently amended) Process The process according to any one of claims claim 9, to 13, characterized in that further comprising the step of, subsequent to step (B), isolating said compound (III) is isolated as a salt by crystallisation from [[the]] said reaction mixture.
- 15. (currently amended) Process The process according to any one of claims claim 9, further comprising the step of, to 14, characterized in that in step (C), adding a base to said reaction mixture to liberate said [[the]] compound (III) is liberated from the salt by addition of a base, preferably by addition of NaOH.
- 16. (currently amended) Process The process according to claim 15, characterized in that wherein step (C) is carried through under heating, and further comprising the step of obtaining with the said compound (III), which is liberated from the salt, being obtained in molten form.
- 17. (currently amended) Process The process according to claim 16, characterized in that the further comprising the step of cooling down said compound (III) present in molten form by freezing is cooled down and, after freezing, is crystallized crystallizing said compound (III) from an aqueous alkaline solution.
- 18. (currently amended) Process A process for the production of a compound (III), starting from a salt of said compound (III), characterized in that comprising the step of liberating and isolating said compound (III) is liberated and isolated from the salt as a free base in molten form.
- 19. (currently amended) Process The process according to claim 18, characterized in that the further comprising the step of cooling down said compound (III) present in molten form to freezing is cooled down and, after freezing, is crystallised subsequently crystallizing said compound (III) from an aqueous alkaline solution.

 20. (currently amended) Process The process for the production of a compound of formula (III) according to claim 18, further comprising the as indicated, characterized in that it contains a step of separating said wherein this compound (III) is separated from the reaction mixture in liquid form.
- 21. (currently amended) Process The process for the production of a compound of formula (III) as indicated, according to claim 20, further comprising characterized in that said process comprises the following steps:
- reducing the aforementioned said compound (I) to compound (III), which gives to provide said compound (III) in salt form; and

- adding a base to said compound (III) to liberate said, whereby compound (III) is liberated from the salt and to separate said compound (III) separates out in liquid form.
- 22. (currently amended) Use of a compound of formula (I), produced according to a process according to any one of claims claim 1 [[to 9]], for the production of producing a compound of formula (III) in a form selected from the group consisting of [[as]] a free base or in the form of and a salt.
- 23. (new) The process according to claim 3, wherein said process comprises using 2-pyrrolidone in an amount of 2 to 4 mol relative to the amount of compound (II).
- 24. (new) The process according to claim 23, wherein said process comprises using 2-pyrrolidone in an amount of 2.5 to 3.5 mol relative to the amount of compound (II).
- 25. (new) The process according to claim 5, wherein said process comprises the step of initially heating said reaction mixture to an initial temperature of 80 to 120 °C and subsequently heating said reaction mixture to a subsequent temperature of 150 to 190 °C.
- 26. (new) The process according to claim 6, wherein said process comprises the step of maintaining said initial temperature for a period of 1 to 1.5 hours and maintaining said subsequent temperature for a period of 2 to 5 hours.
- 27. (new) The process according to claim 7, comprising the step of maintaining said reaction mixture seeded with said seed crystals at a temperature of at least 25 °C for a period of 50 to 100 hours to enable crystallisation.
- 28. (new) The process according to claim 8, wherein said crystallisation is carried through at a temperature of 40 to 60 °C.
- 29. (new) The process according to claim 15, wherein said base is NaOH.
- 30. (new) Use of a compound of formula (I), produced according to a process according to claim 9, for producing a compound of formula (III) in a form selected from the group consisting of a free base and a salt.